Amendments to the Specification:

At page 1, after the title and before the heading "Background," please insert the following

paragraph:

-- This application is a continuation of U.S. Patent Application Serial No. 09/716,507,

filed November 20, 2000; which is a continuation of U.S. Patent Application Serial

No. 09/244,568, filed February 4, 1999, now Patent No. 6,307,042; which in turn is a

continuation of U.S. Patent Application Serial No. 08/634,053, filed April 17, 1996, now Patent

No. 5,959,098; all of which are hereby incorporated herein by reference.--

Please replace the paragraph beginning at page 28, line 17 with the following paragraph:

-- A schematic illustration of one embodiment of an integrated reactor system is shown in

Figure 3C 4C. The device includes an automated peptide synthesizer 401. The automated

peptide synthesizer is a device which flows selected reagents through a flow cell 402 under the

direction of a computer 404. In a preferred embodiment the synthesizer is an ABI Peptide

Synthesizer, model no. 431A. The computer may be selected from a wide variety of computers

or discrete logic including for, for example, an IBM PC-AT or similar computer linked with

appropriate internal control systems in the peptide synthesizer. The PC is provided with signals

from the ABI computer indicative of, for example, the beginning of a photolysis cycle. One can

also modify the synthesizer with a board that links the contacts of relays in the computer in

parallel with the switches to the keyboard of the control panel of the synthesizer to eliminate

some of the keystrokes that would otherwise be required to operate the synthesizer.--

Cont. of USSN 09/716,507 Express Mail Receipt No. EV 323344325 US 2

Please replace the paragraph beginning at page 55, line 28 with the following paragraph:

-- Figures 8A and 8B 9A and 9B illustrate the contrast difference between back-side

exposure synthesis and front-side exposure synthesis, respectively. Figure 9A shows a

fluorescent scan of a substrate having fluorescent groups coupled directly to the surface of the

substrate using photolithographic techniques, with a mask having 50 µm and 100 µm feature

sizes where the activating light was shown through the back-side of the substrate. Figure 9B

shows the same synthesis where the activation light was directed at the front side of the

substrate. The definition of the individual features is greatly enhanced using this front-side

photolysis.--

[REMAINDER OF PAGE INTENTIONALLY BLANK]

3

Cont. of USSN 09/716,507 Express Mail Receipt No. EV 323344325 US